

Pulmonary Pathology for the Boards



Robert F. Padera, M.D., Ph.D.

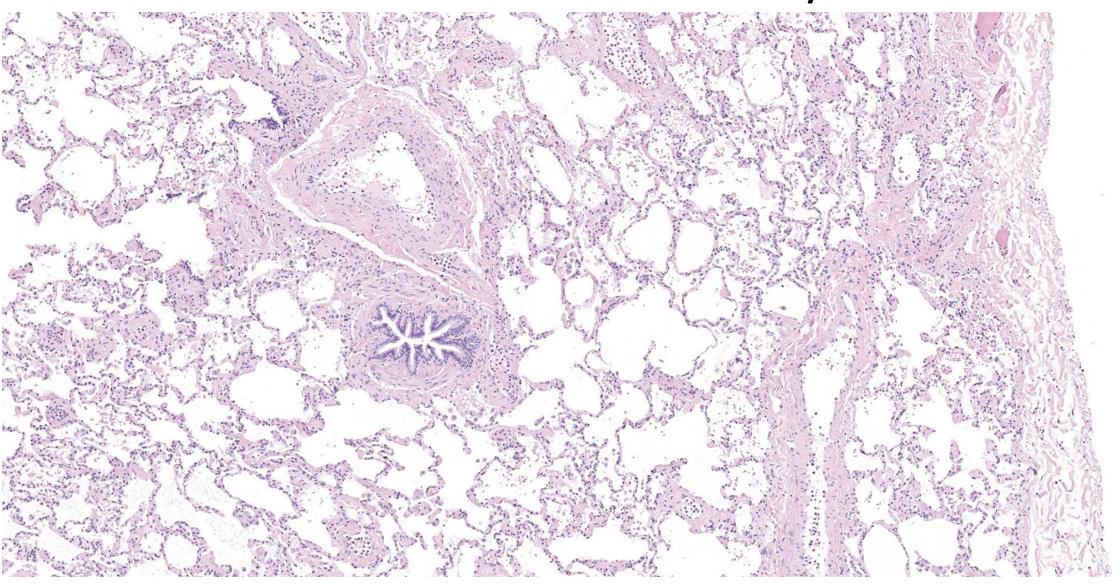
Associate Professor of Pathology, Harvard Medical School
Pathologist, Brigham and Women's Hospital
Boston, Massachusetts
rpadera@bwh.harvard.edu

Conflict of Interest Disclosure: None

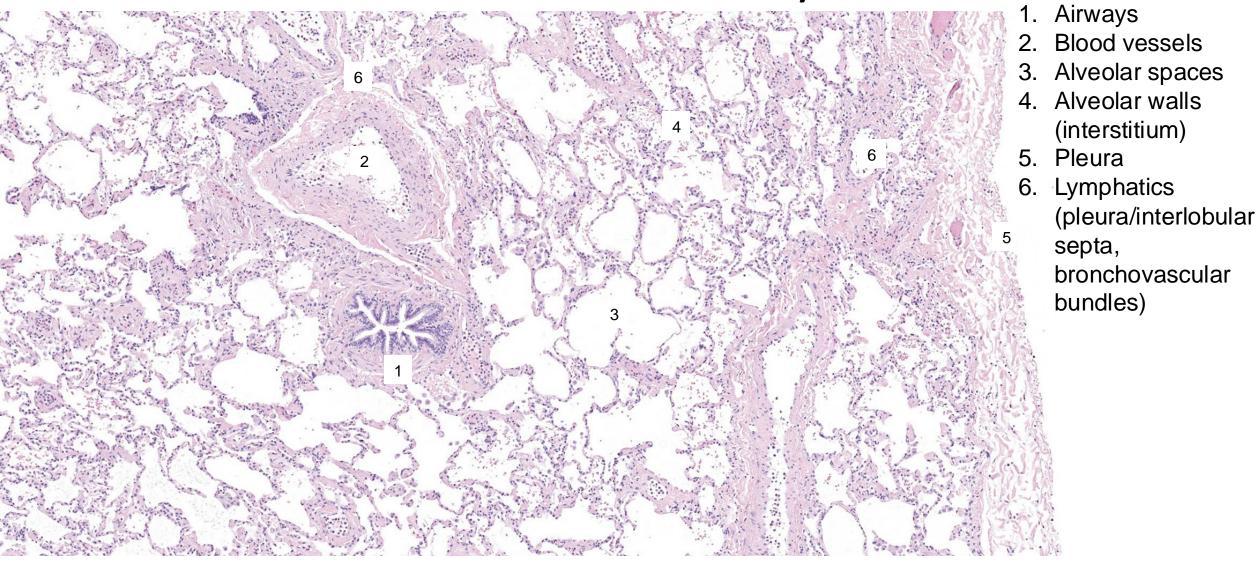
Goals

- To teach you everything about pulmonary pathology
- To review commonly encountered non-neoplastic and neoplastic pulmonary pathology that might be encountered on the boards

Normal Anatomy



Normal Anatomy



Outline

- Interstitial lung disease
 - Infection
 - Malignancy
 - Pathology special stains and buzzwords

Interstitial Lung Disease

- Usual interstitial pneumonia (UIP)
- Nonspecific interstitial pneumonia (NSIP)
- Acute interstitial pneumonia (AIP)
- Cryptogenic organizing pneumonia (COP)
- Hypersensitivity pneumonitis (HP)
- Sarcoidosis

Not a comprehensive list

ILD – General Comments

- Clinical-radiologic-pathologic correlation
 - Multidisciplinary meeting
- Histologic patterns of disease
- Significant infidelity
 - One etiology can give rise to several patterns
 - Same pattern can be present in multiple diseases
- Surgical vs. transbronchial biopsy

Usual Interstitial Pneumonia - Gross

Honeycombing, most severe/earliest in lower zones, subpleural areas



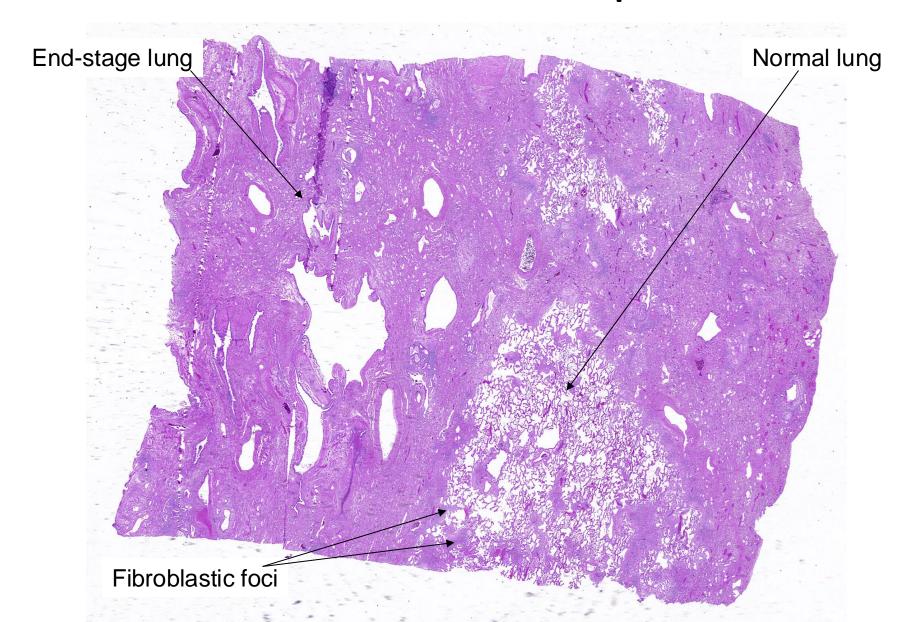


Pleural surface

UIP - Microscopic

- Surgical biopsies multiple lobes/areas
- Temporal and spatial heterogeneity patchy
 - End-stage ("honeycomb") lung
 - Fibroblastic foci
 - Normal lung
 - Minimal active inflammation (usually)
- Architectural distortion fibrosis
- Absence of other specific findings (e.g., granulomas, vasculitis, aspirated material)

UIP - Microscopic

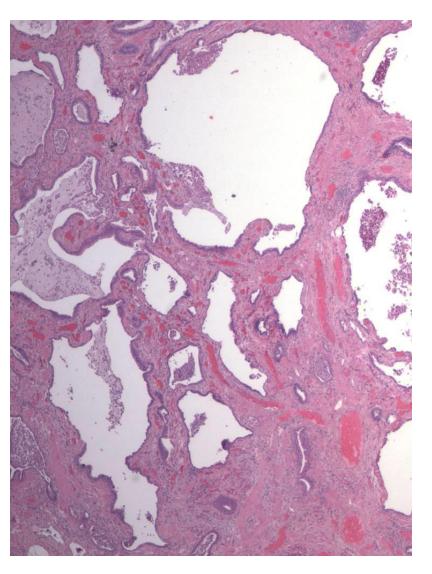


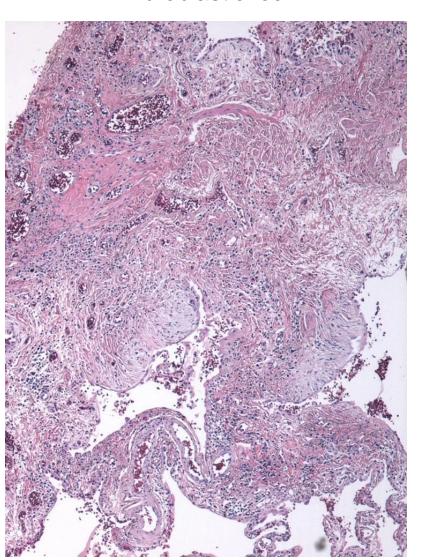
UIP - Microscopic

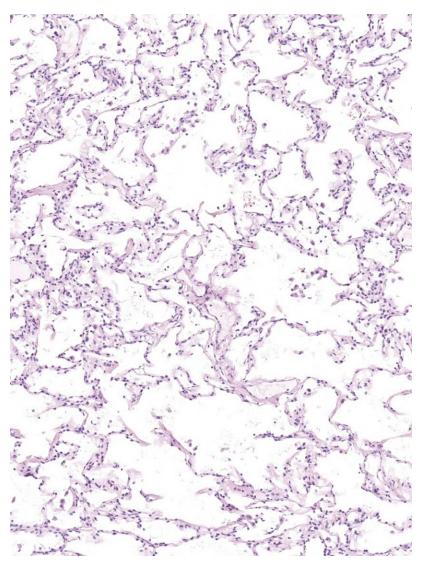
End-stage lung

Fibroblastic foci

Normal lung





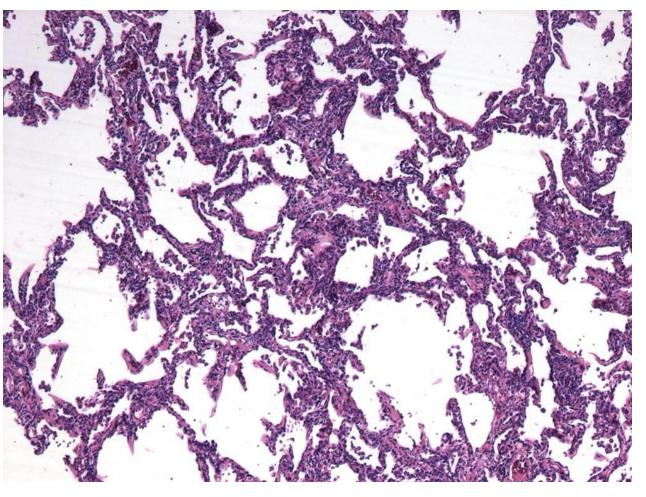


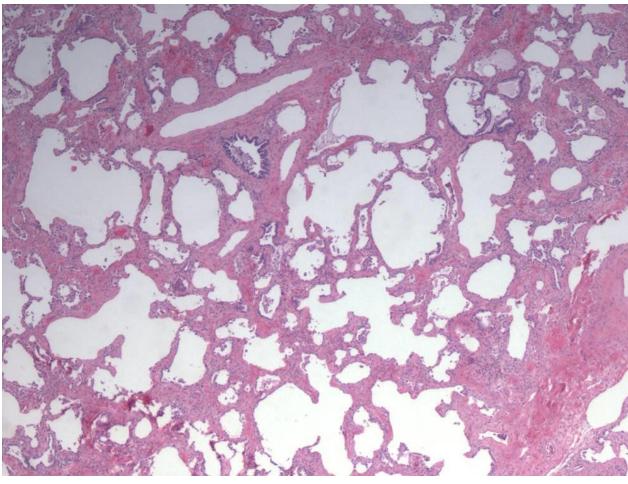
Nonspecific Interstitial Pneumonia

- Temporal and spatial homogeneity
- Cellular NSIP chronic inflammation
- Fibrotic NSIP fibrosis, less inflammation
- Architecture maintained

Nonspecific Interstitial Pneumonia

Cellular NSIP Fibrotic NSIP

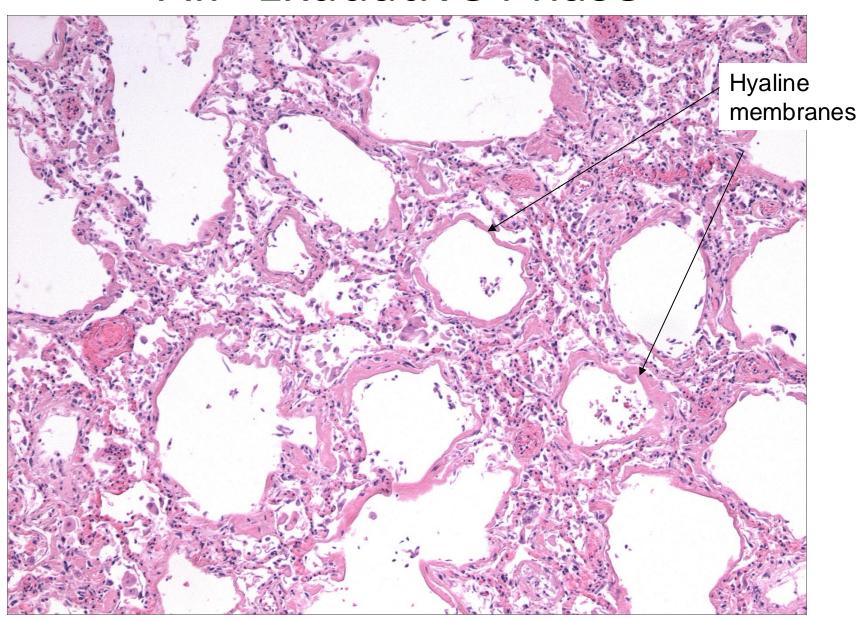




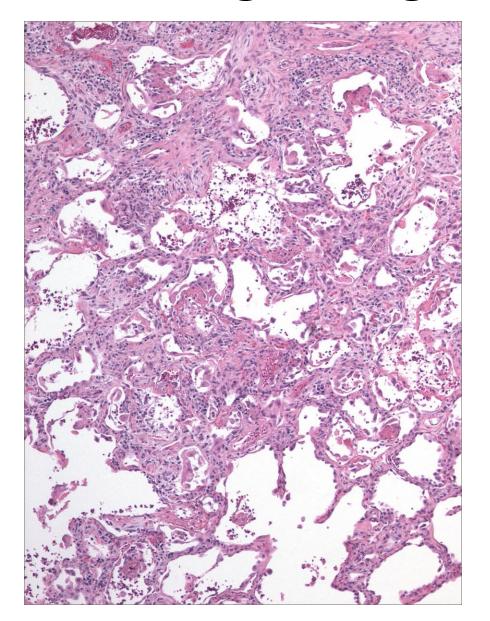
Acute Interstitial Pneumonia

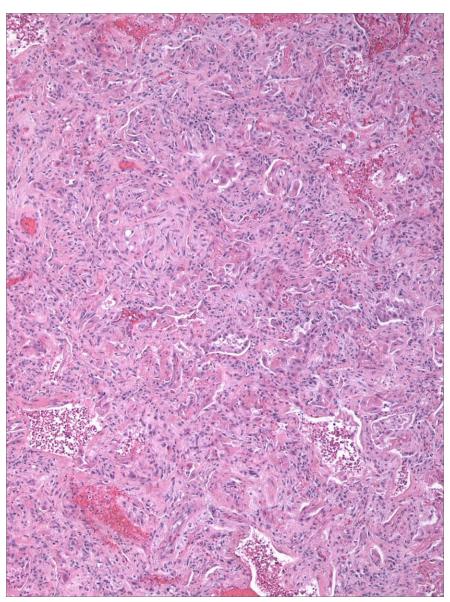
- "Idiopathic" acute respiratory distress syndrome – no underlying cause (e.g., sepsis, toxic inhalation, pancreatitis)
- Pathologic findings acute/exudative, organizing, fibrotic <u>diffuse alveolar damage</u> without morphologic clues as to etiology

AIP-Exudative Phase



AIP- Organizing and Fibrotic Phase

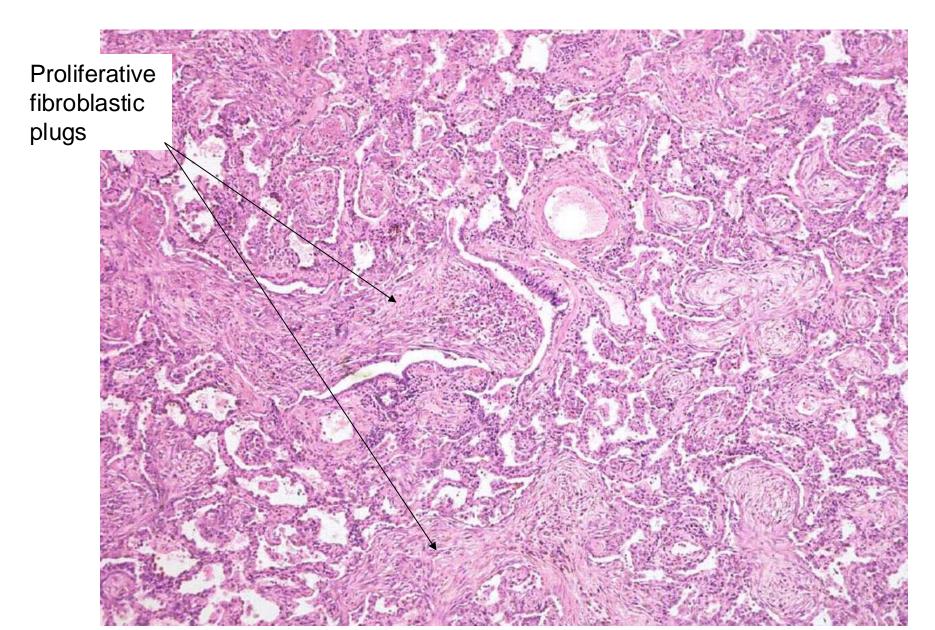




Cryptogenic Organizing Pneumonia

- "Idiopathic" bronchiolitis obliterans and organizing pneumonia
- Proliferative fibroblastic plugs within respiratory bronchioles, alveolar ducts, alveolar spaces
- Variable interstitial chronic inflammation
- Temporally homogeneous with preservation of underlying lung architecture
- Histologic pattern can be seen in various clinical scenarios and as part of other specific pathologic entities (e.g., hypersensitivity pneumonitis, at periphery of malignancies, resolving infection)

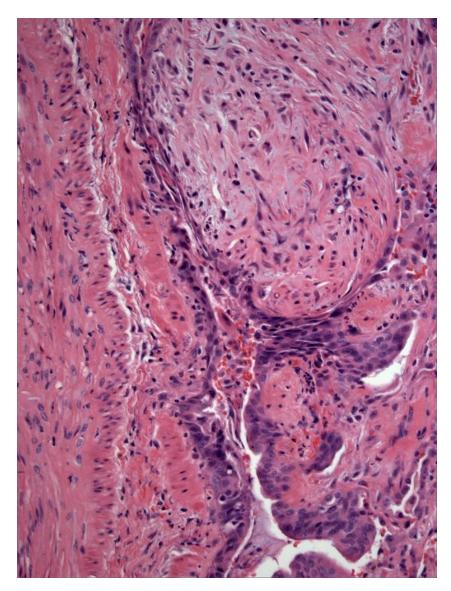
Cryptogenic Organizing Pneumonia

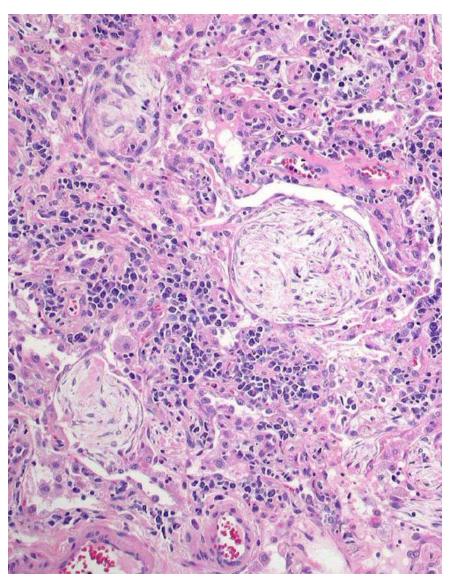


Cryptogenic Organizing Pneumonia

Bronchiolitis obliterans

Organizing pneumonia

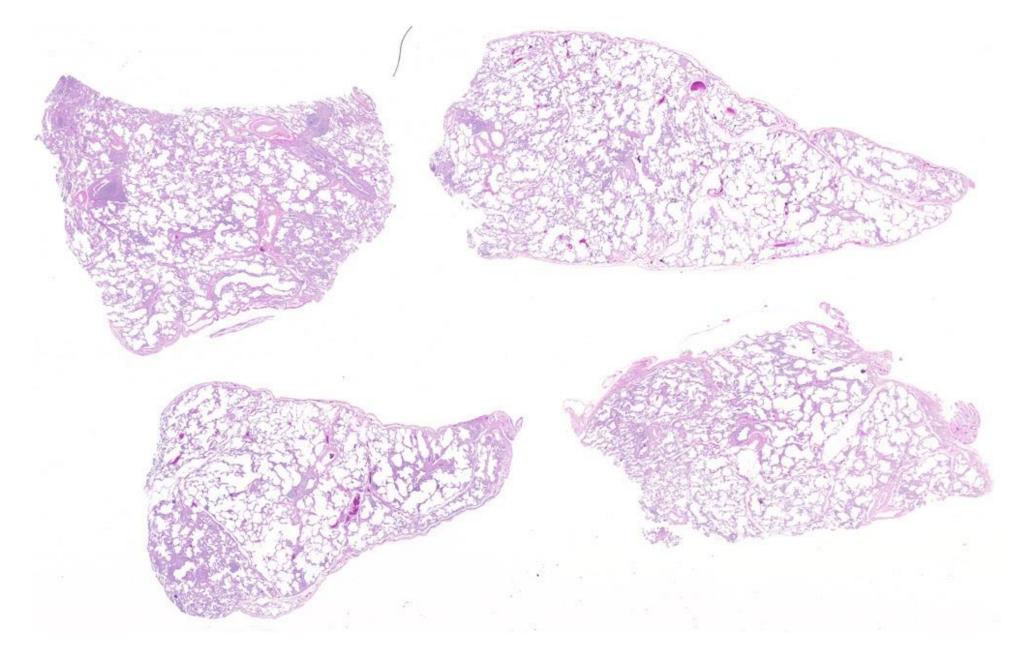




Hypersensitivity Pneumonitis

- Patchy interstitial pneumonitis with lymphocytes and plasma cells, accentuated around small airways
- Loosely formed non-necrotizing granulomas
- Foci of organizing pneumonia

Hypersensitivity Pneumonitis

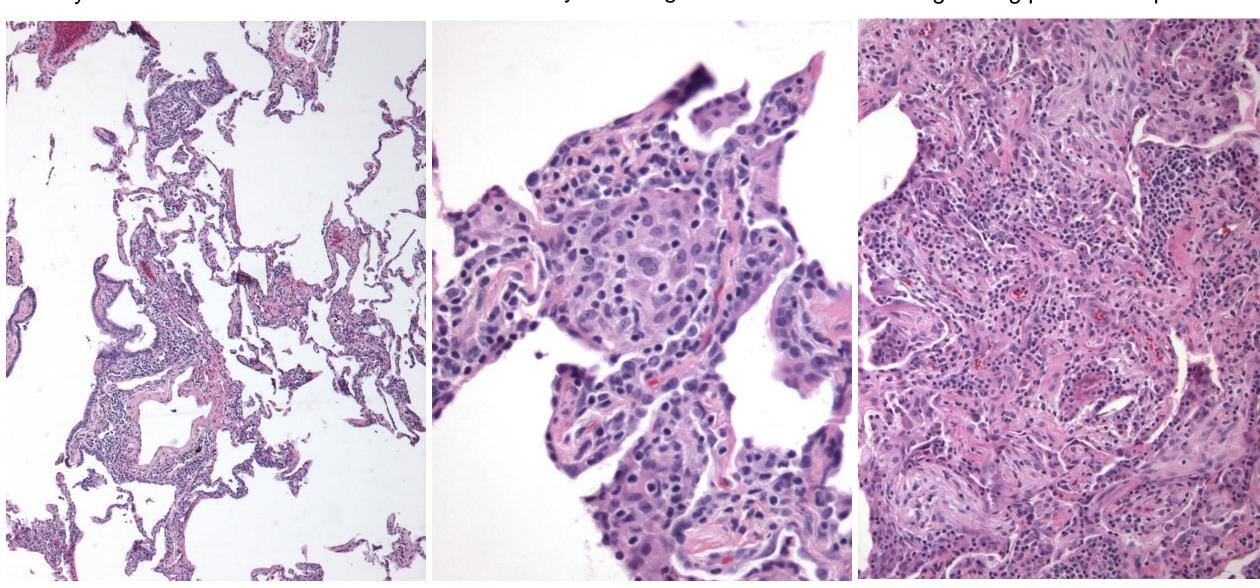


Hypersensitivity Pneumonitis

Airway-centered chronic inflammation

Loosely formed granulomas

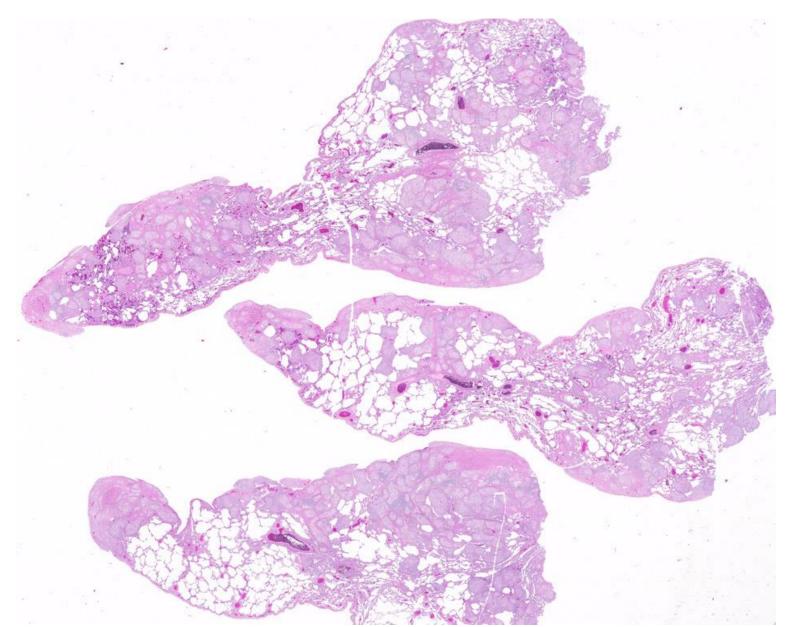
Organizing pneumonia pattern



Sarcoidosis

- Well-formed non-necrotizing granulomas
- Often with surrounding fibrosis and variable chronic inflammation
- Lymphatic distribution peribronchiolar and septal/pleural
- Transbronchial biopsy can be diagnostic multiple samples, multiple levels

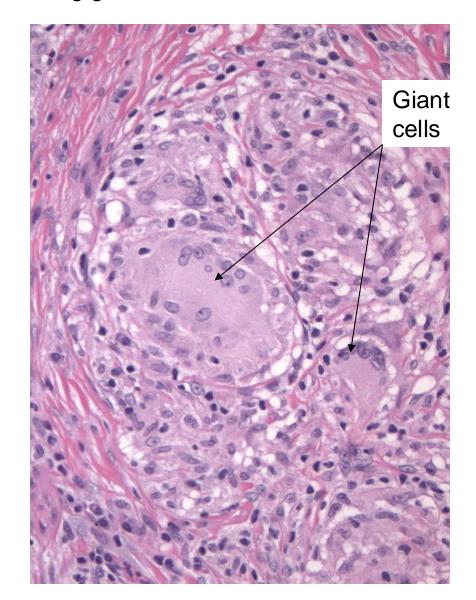
Sarcoidosis



Sarcoidosis

Well-formed non-necrotizing granulomas





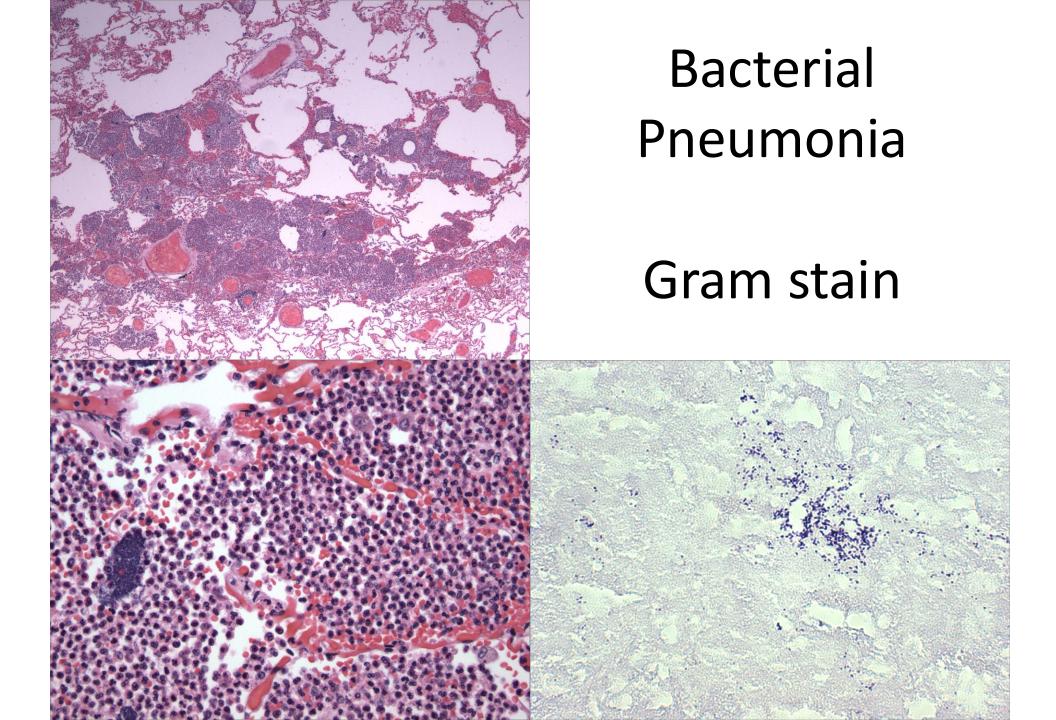
Outline

Interstitial lung disease

- Infection
 - Malignancy
 - Pathology special stains and buzzwords

Bacterial Pneumonia

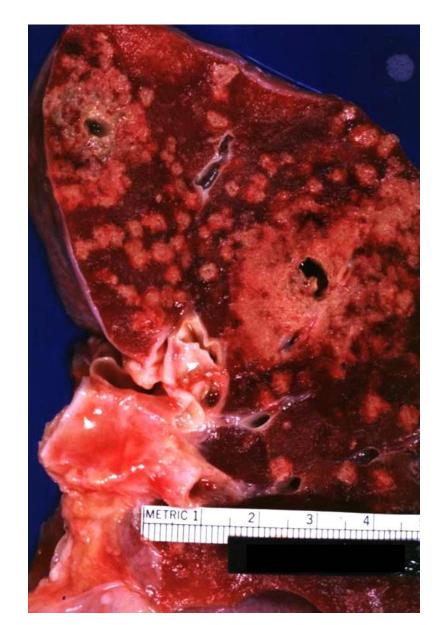
- Acute inflammation neutrophils, fibrin, edema, bacteria in alveolar spaces
- Gram stain can highlight bacterial organisms in tissue sections and smears
- Progression to organizing pneumonia that resembles BOOP/COP discussed prior

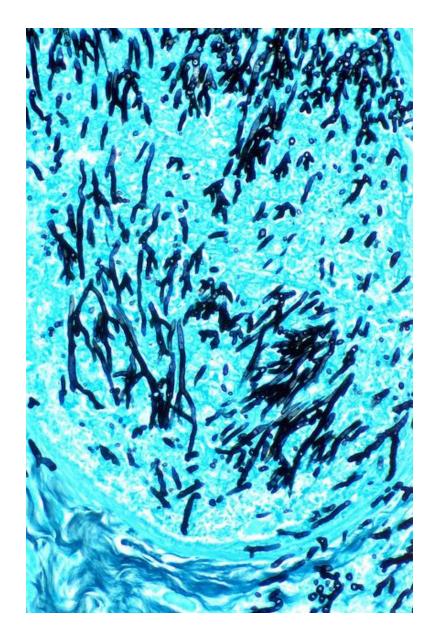


Fungal Pneumonia

- Many fungi will elicit a granulomatous reaction
 may be necrotizing or not
- Special stains to identify organisms include methenamine silver stain (MSS), Gomori methenamine stain (GMS) and periodic acid-Schiff (PAS) stain
- Pneumocystis jiroveci produces a cell-poor, proteinaceous alveolar infiltrate

Aspergillus Pneumonia – MSS stain

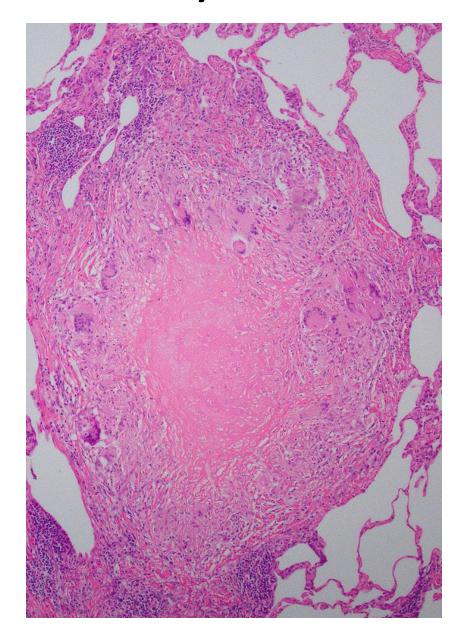


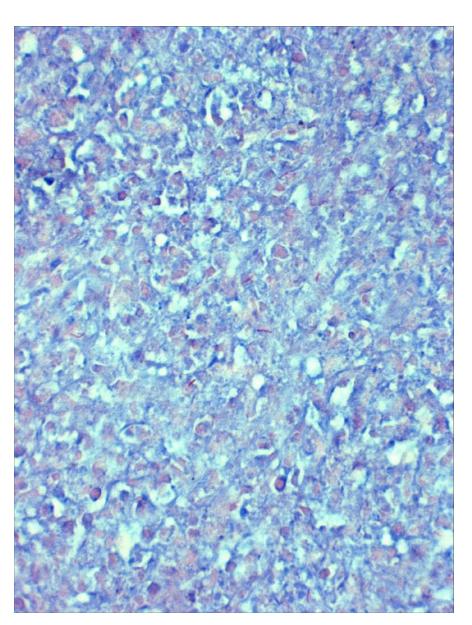


Mycobacterial Infection

- Most mycobacteria will elicit a granulomatous reaction – may be necrotizing or not
- Special stains to identify organisms include acid-fast bacilli (AFB) stain and modifications thereof (e.g., Nocardia stain)
- Infiltrates of foamy macrophages may contain non-tuberculous mycobacteria in immunocompromised patients

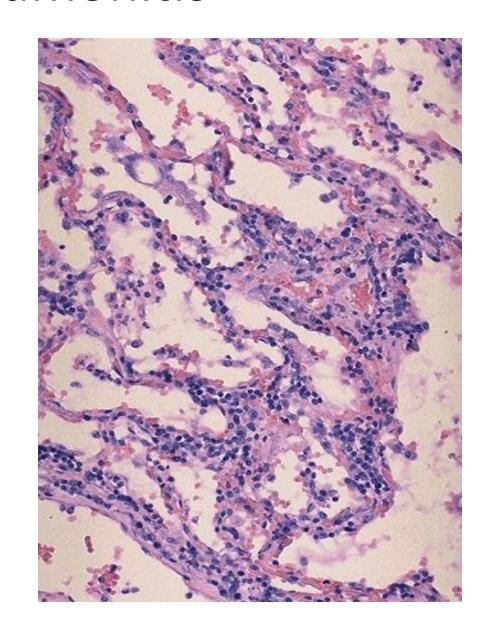
Mycobacterial Infection - AFB





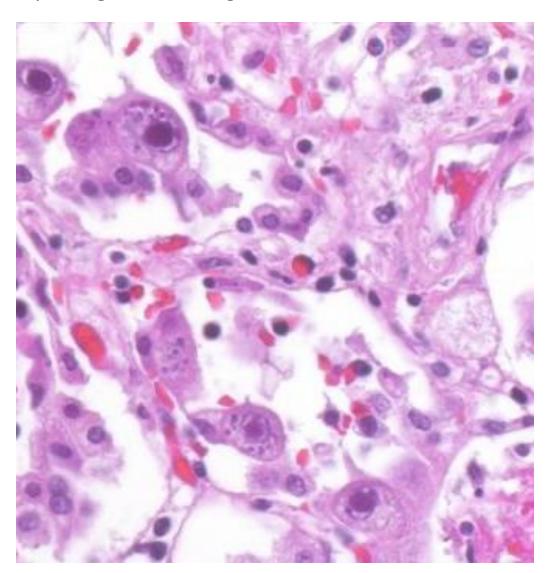
Viral Pneumonitis

- Milder injury pattern for many respiratory viral infections (RSV, parainfluenza, adenovirus, metapneumovirus, et al.)
- Variable interstitial mononuclear cell infiltrates, edema
- Severe viral infection may produce DAD/ARDS pattern (SARS-CoV-2)

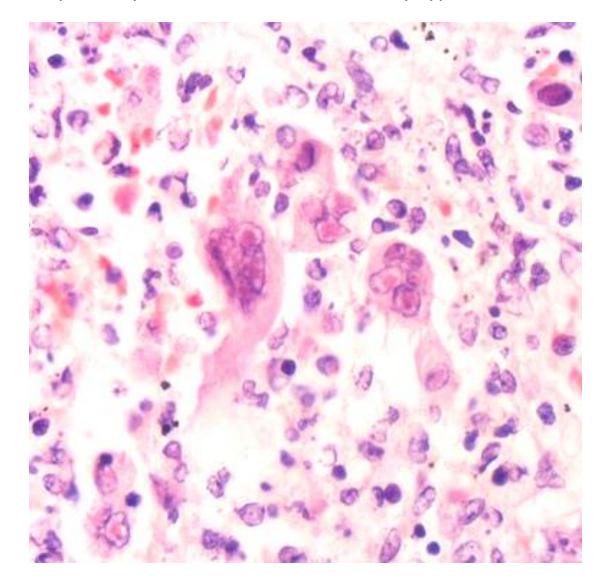


Diagnostic Viral Inclusions

Cytomegalovirus: Large cells, red nuclear inclusions



Herpes simplex: Giant cells with Cowdry type A inclusions



Outline

- Interstitial lung disease
- Infection
- Malignancy
 - Pathology special stains and buzzwords

Common malignancies

- Adenocarcinoma
- Squamous cell carcinoma
- Carcinoid/atypical carcinoid
- Small cell carcinoma
- (Metastasis)

Adenocarcinoma

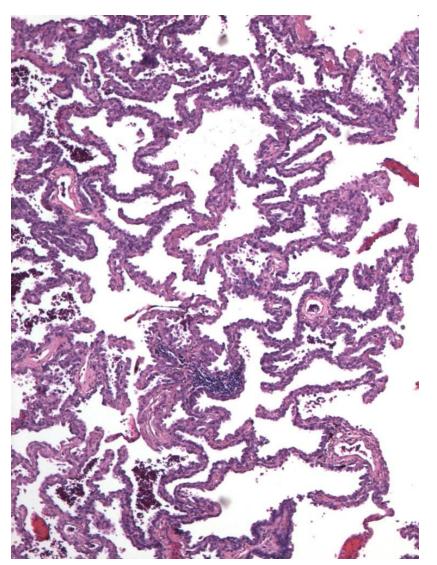
- Atypical adenomatous hyperplasia
- Adenocarcinoma in situ (formerly non-mucinous BAC)
- Minimally invasive adenocarcinoma
- Invasive adenocarcinoma, predominantly...
 - Lepidic, acinar, papillary, micropapillary or solid
- Variants
 - Invasive mucinous adenocarcinoma (formerly mucinous BAC)
- Genetic studies EGFR, EML4-ALK, ROS1, KRAS

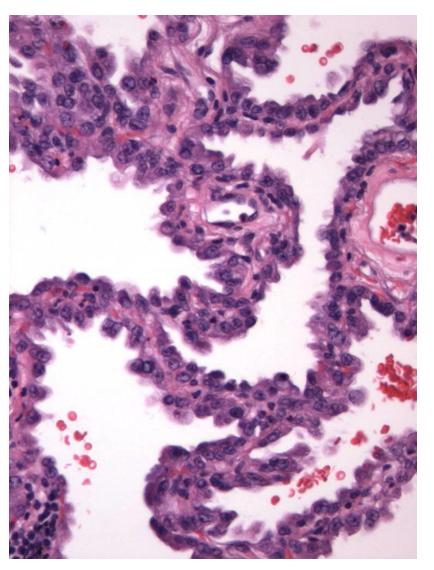
Adenocarcinoma

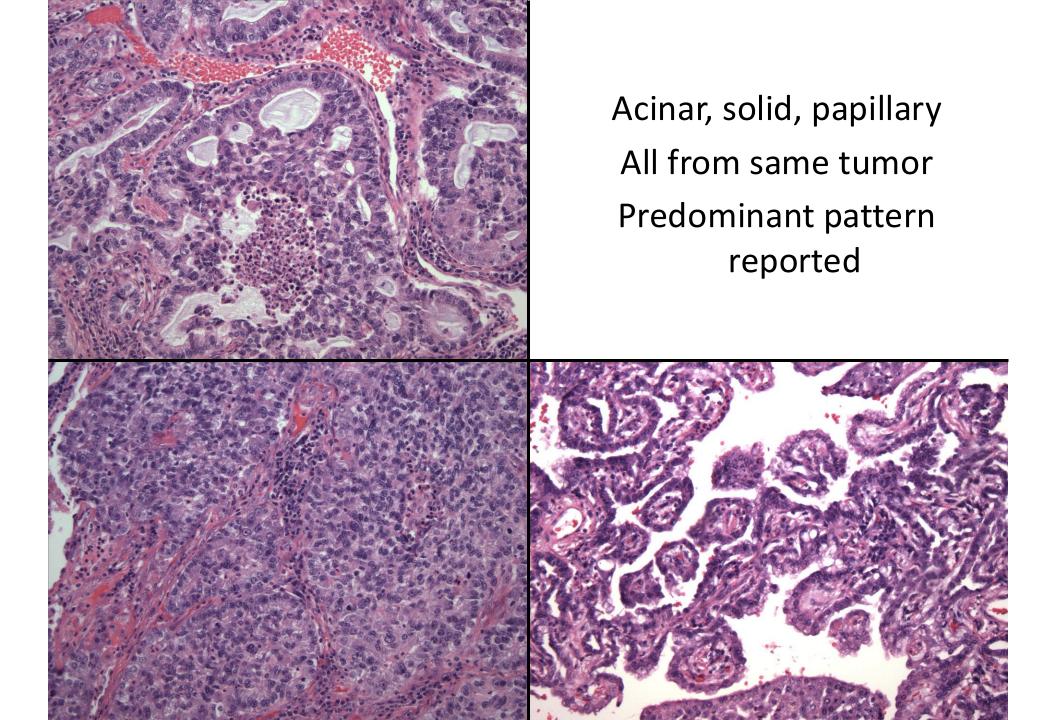


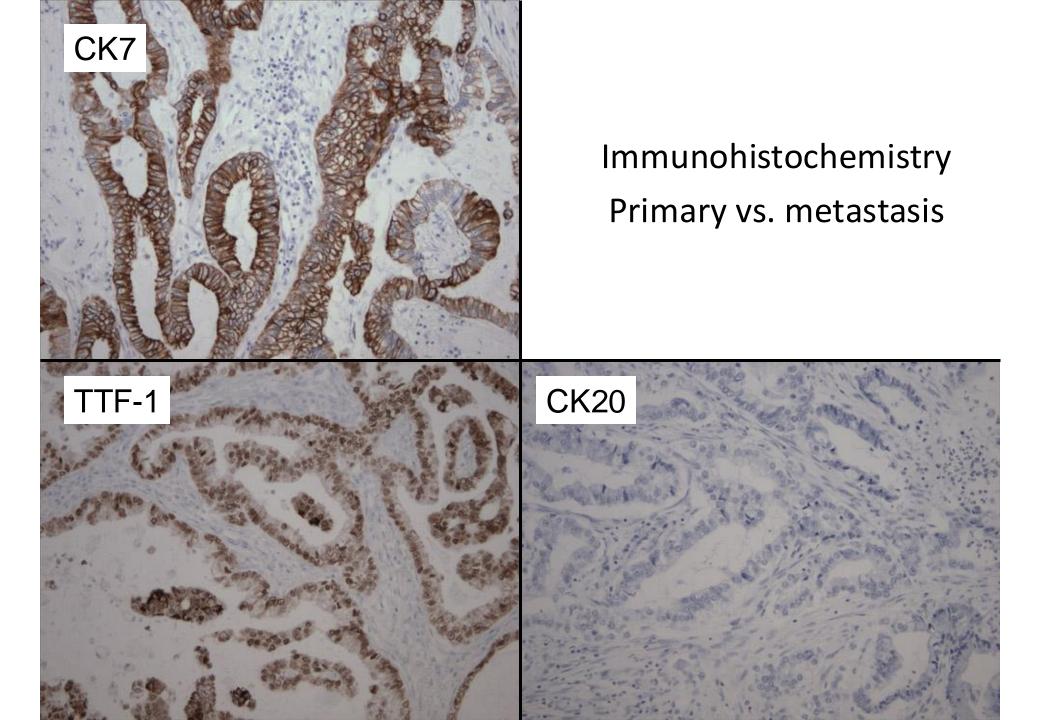
Adenocarcinoma In Situ

Growth along pre-existing alveolar walls with no stromal invasion

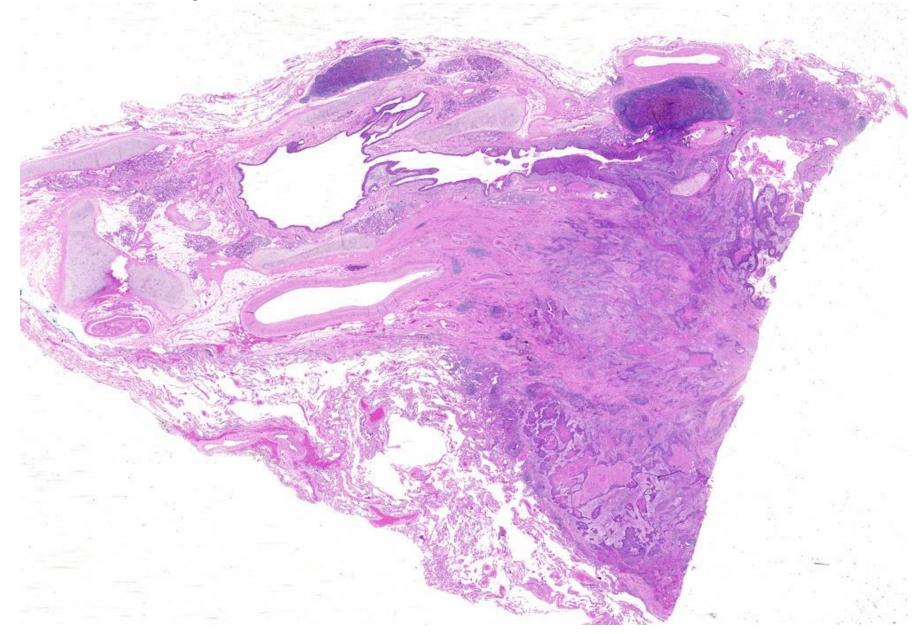




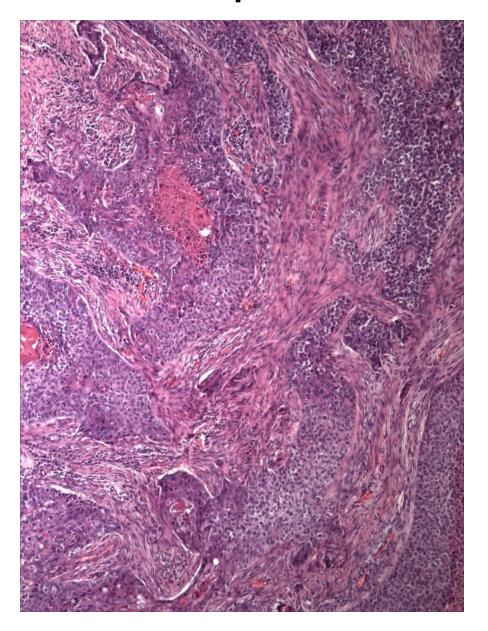


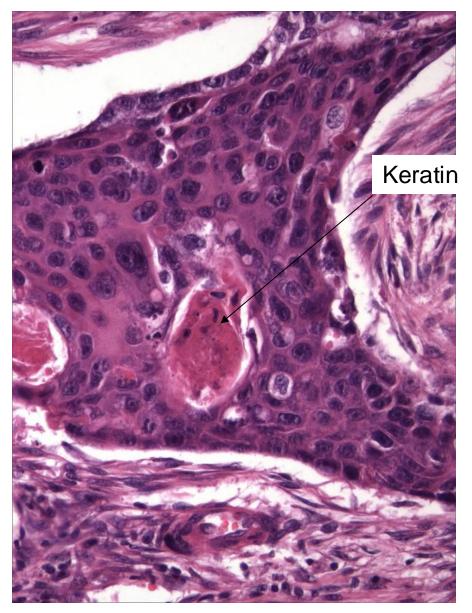


Squamous Cell Carcinoma

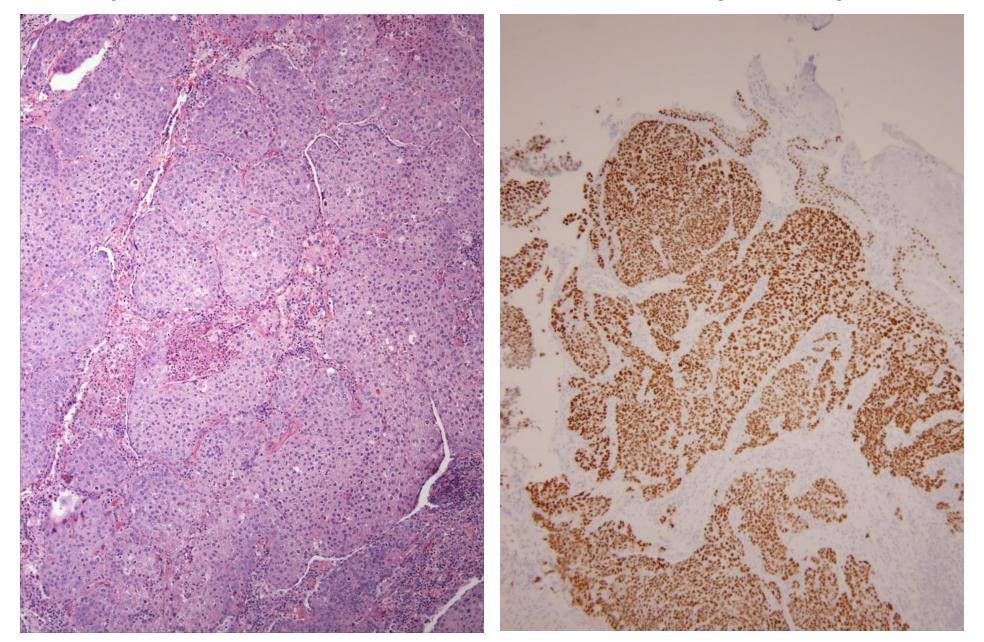


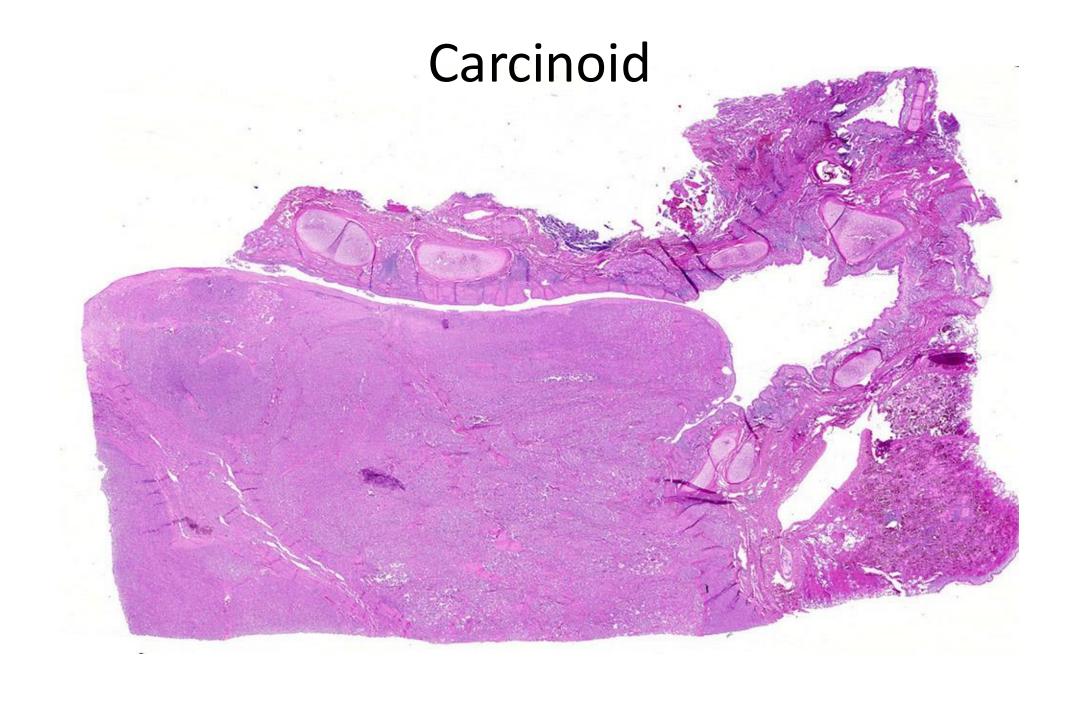
Squamous Cell – Invasive



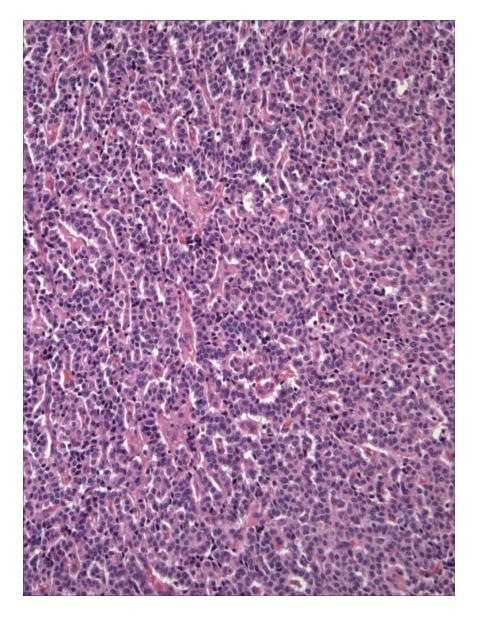


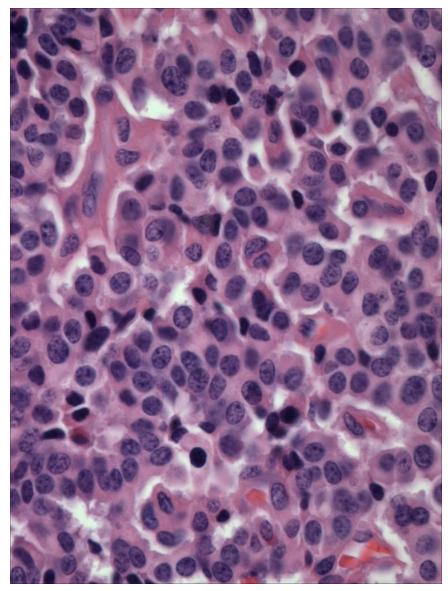
Squamous Cell – Invasive – p40, p63





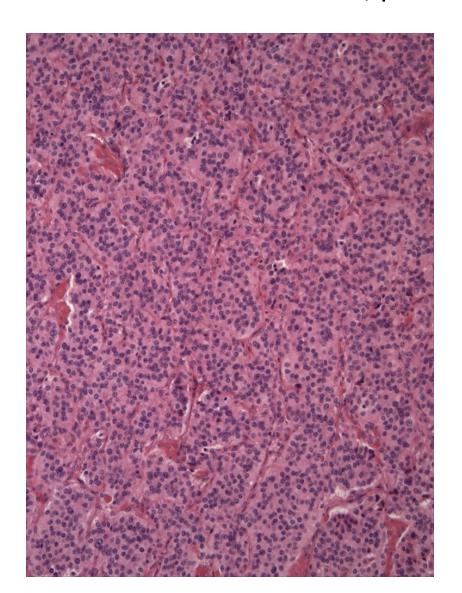
Carcinoid

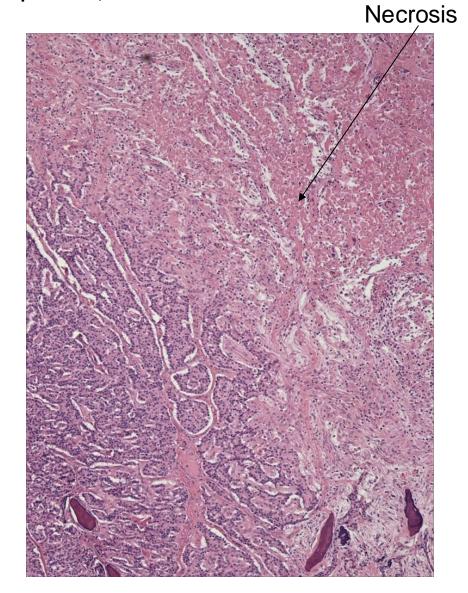




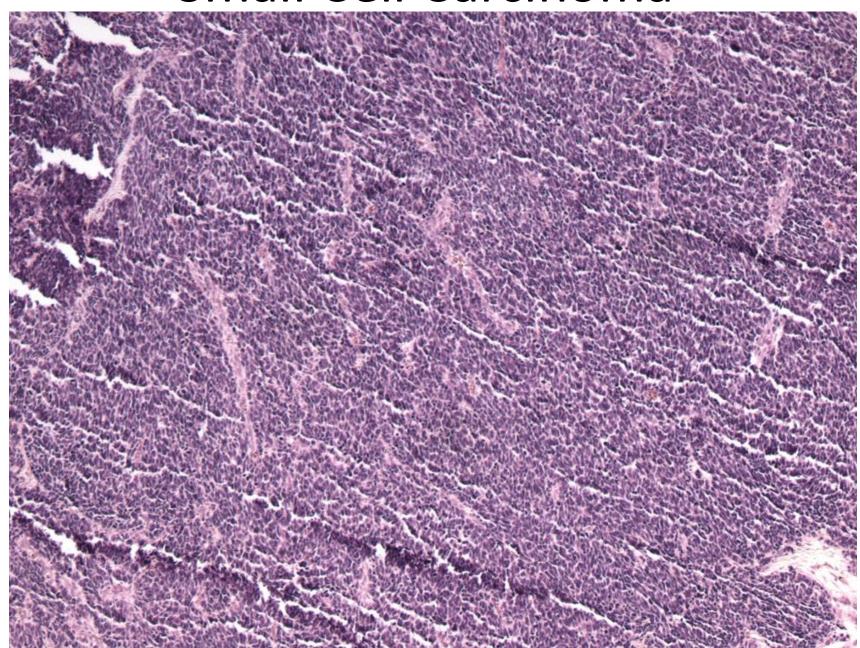
Atypical Carcinoid

Increased mitosis, pleomorphism, areas of necrosis





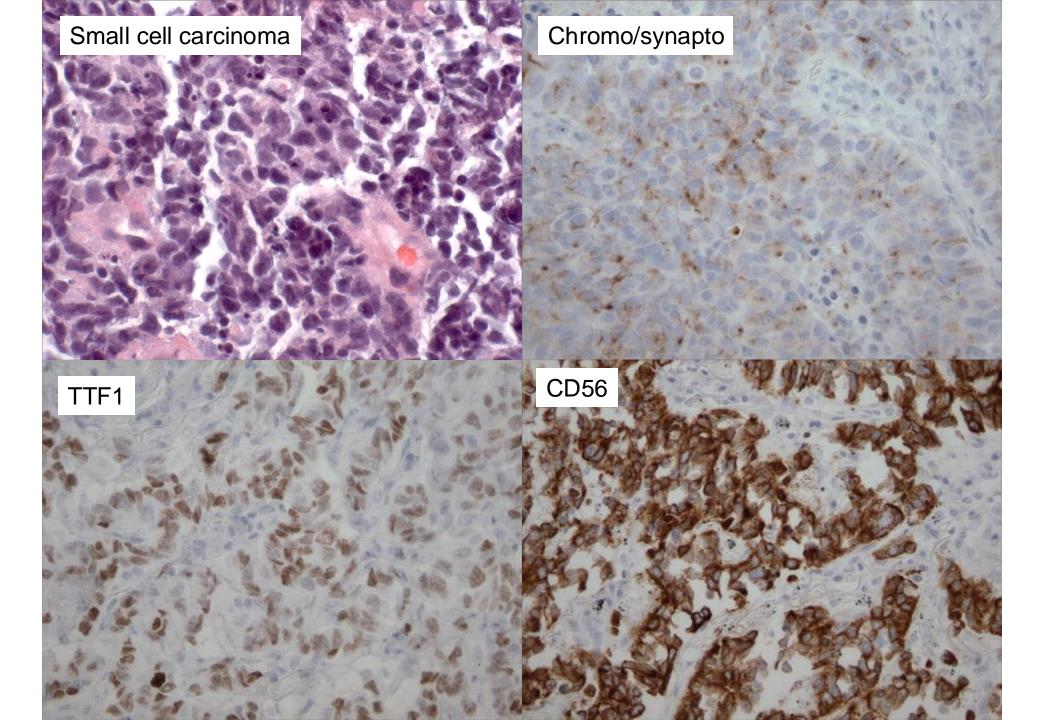
Small Cell Carcinoma



Small Cell Carcinoma

Hyperchromasia, nuclear molding, mitoses, single cell necrosis

Necrosis Mitoses



Outline

- Interstitial lung disease
- Infection
- Malignancy

Pathology special stains and buzzwords

Neoplastic Special Stains

- Pulmonary adenocarcinoma
 - TTF-1, napsin, CK7, mucicarmine positive
 - p40, p63 negative
- Squamous cell carcinoma
 - p40, p63, CK5/6 positive
 - TTF-1, napsin negative
- Small cell carcinoma
 - Keratin, TTF-1, CD56, synaptophysin (50%),
 chromogranin (50%) positive
- Immunohistochemistry for ALK, ROS1, PD-L1 (also some EGFR and BRAF mutants)

Non-neoplastic Special Stains

- Trichrome stain fibrosis
- Congo red stain, apple green birefringence amyloid
- AFB stain mycobacteria, Nocardia
- MSS, GMS fungus
- PAS fungus; also for pulmonary alveolar proteinosis
- Gram stain bacteria

Buzzwords

- Spatial/temporal heterogeneity, fibroblastic foci UIP
- Hyaline membranes DAD/AIP
- Loosely formed granulomas HP
- Well-formed granulomas sarcoid
- Necrotizing granulomas TB, Fungus, GPA
- Nuclear molding, single cell necrosis small cell carcinoma
- Central tumor in smoker squamous cell or small cell carcinoma
- Central tumor in nonsmoker carcinoid
- Peripheral tumor, lepidic growth adenocarcinoma



Pulmonary Pathology for the Boards



Robert F. Padera, M.D., Ph.D.

Associate Professor of Pathology, Harvard Medical School
Pathologist, Brigham and Women's Hospital
Boston, Massachusetts
rpadera@bwh.harvard.edu